This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

(Currently Amended) A compound of the formula

$$\underset{R^1}{\overset{\bullet}{\bigcap}} \underset{R^2}{\overset{\bullet}{\bigcap}}$$

in which

R¹ is C₁-C₈-alkyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl or C₃-C₈-cycloalkyl, where C₁-C₈-alkyl is optionally substituted by oxo, and

where C_1 - C_8 -alkyl, C_2 - C_6 -alkenyl, C_2 - C_6 -alkynyl and C_3 - C_8 -cycloalkyl are optionally substituted by up to 3 radicals independently of one another selected from the group of C_1 - C_6 -alkyl, C_1 - C_6 -alkoxy, hydroxycarbonyl, cyano, amino, nitro, hydroxy, C_1 - C_6 -alkylamino, halogen, trifluoromethyl, trifluoromethoxy.

 $C_6\text{-}C_{10}\text{-}arylcarbonylamino, $C_1\text{-}C_6\text{-}alkylcarbonylamino, $C_1\text{-}C_6\text{-}alkylamino-carbonyl, $C_1\text{-}C_6\text{-}alkoxycarbonyl, $C_6\text{-}C_{10}\text{-}arylaminocarbonyl, heteroarylamino-carbonyl, heteroarylcarbonylamino, $C_1\text{-}C_6\text{-}alkylsulphonylamino, $C_1\text{-}C_6\text{-}alkylsulphonyl,}$$

C1-C6-alkylthio,

where

$$\begin{split} &C_1\text{--}C_6\text{-}alkyl, C_1\text{--}C_6\text{-}alkoxy, C_1\text{--}C_6\text{-}alkylamino, C_6\text{--}C_{10\text{-}}arylcarbonylamino, C_1\text{--}\\ &C_6\text{-}alkylcarbonylamino, C_1\text{--}C_6\text{-}alkylaminocarbonyl, C_1\text{--}C_6\text{--}alkoxy-carbonyl,} \end{split}$$

C6-C10-arylaminocarbonyl, heteroarylaminocarbonyl,

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heteroarylcarbonylamino, C_1 - C_6 -alkylsulphonylamino, C_1 - C_6 -alkylsulphonyl and C_1 - C_6 -alkylthio are optionally substituted by one to three radicals independently of one another selected from the group of hydroxy, cyano, halogen, trifluoromethyl, trifluoromethoxy, hydroxy-carbonyl and a group of the formula $-NR^3R^4$,

where

R³ and R⁴ are independently of one another hydrogen or C₁-C₆-alkyl,

or

R³ and R⁴ together with the nitrogen atom to which they are bonded are 5to 8-membered heterocyclyl,

R² is phenyl or heteroaryl, where phenyl is substituted by 1 to 3 radicals and heteroaryl is optionally substituted by 1 to 3 radicals in each case independently of one another selected from the group of C₁-C₆-alkyl, C₁-C₆-alkoxy, hydroxycarbonyl, cyano, trifluoromethyl, trifluoromethoxy, amino, nitro, hydroxy, C₁-C₆-alkylamino, halogen, C₆-C₁₀-arylarbonylamino, C₁-C₆-alkylaminocarbonyl, C₁-C₆-alkoxycarbonyl, C₆-C₁₀-arylaminocarbonyl, heteroarylaminocarbonyl, heteroarylamino, C₁-C₆-alkylsulphonyl and C₁-C₆-alkylthio.

where C₁-C₆-alkyl, C₁-C₆-alkoxy, C₁-C₆-alkylamino, C₆-C₁₀-arylcarbonylamino, C₁-C₆-alkylcarbonylamino, C₁-C₆-alkylaminocarbonyl, C₁-C₆alkoxycarbonyl, C₆-C₁₀-arylaminocarbonyl, heteroarylaminocarbonyl, heteroarylcarbonylamino, C₁-C₆-alkylsulphonylamino, C₁-C₆alkylsulphonyl and C₁-C₆-alkyltnio are optionally substituted by one to three radicals independently of one another selected from the group of hydroxy, cyano, halogen, trifluoromethyl, trifluoromethoxy, hydroxycarbonyl and a group of the formula -NR³R⁴.

where

R3 and R4 have the meanings indicated above.

2. (Currently Amended) The compound of Claim claim 1, where

 \mathbb{R}^1 is C1-C8-alkyl, C2-C6-alkenyl, C2-C6-alkynyl or C3-C8-cycloalkyl, which are optionally substituted by up to 3 radicals independently of one another selected from the group of C1-C6-alkyl, C1-C6-alkoxy, hydroxycarbonyl, cyano, amino, nitro, hydroxy, C1-C6-alkylamino, halogen, C6-C10-arylcarbonylamino, C1-C6-alkylcarbonylamino, C1-C6-alkylaminocarbonyl, C1-C6alkoxycarbonyl, C6-C10-arylaminocarbonyl, heteroarylaminocarbonyl, heteroarylcarbonylamino, C1-C6-alkylsulphonylamino, C1-C6-alkylsulphonyl and C1-C6-alkylthio,

where C1-C6-alkyl, C1-C6-alkoxy, C1-C6-alkylamino, C6-C10-arylcarbonylamino, C1-C6-alkylcarbonylamino, C1-C6-alkylaminocarbonyl, C1-C6-alkoxycarbonyl.

C6-C10-arylaminocarbonyl, heteroarylaminocarbonyl, heteroarylcarbonylamino,

C1-C6-alkylsulphonylamino, C1-C6-alkylsulphonyl and C1-C6-alkylthio are optionally substituted by a radical selected from the group of hydroxy, cyano, halogen, hydroxycarbonyl and a group of the formula -NR3R4.

where

R3 and R4 are independently of one another hydrogen or C₁-C₆-alkyl,

or

 \mathbb{R}^2

R³ and R⁴ together with the nitrogen atom to which they are bonded are 5to 8-membered heterocyclyl,

heteroaryl is optionally substituted by 1 to 3 radicals in each case independently of one another selected from the group of C₁-C₆-alkyl, C₁-C₆alkoxy, hydroxycarbonyl, cyano, trifluoromethyl, amino, nitro, hydroxy, C1-5

is phenyl or heteroaryl, where phenyl is substituted by 1 to 3 radicals and

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 C_6 -alkylamino, halogen, C_6 - C_{10} -arylcarbonylamino, C_1 - C_6 -alkylcarbonylamino, C_1 - C_6 -alkylaminocarbonyl, C_1 - C_6 -alkoxycarbonyl, C_6 - C_{10} -arylaminocarbonyl, heteroarylcarbonylamino, C_1 - C_6 -alkylsulphonylamino, C_1 - C_6 -

where C₁-C₆-alkyl, C₁-C₆-alkoxy, C₁-C₆-alkylamino, C₆-C₁₀-arylcarbonylamino, C₁-C₆-alkylcarbonylamino, C₁-C₆-alkylaminocarbonyl, C₁-C₆alkoxycarbonyl, C₆-C₁₀-arylaminocarbonyl, heteroarylaminocarbonyl, heteroarylcarbonylamino, C₁-C₆-alkylsulphonylamino, C₁-C₆alkylsulphonyl and C₁-C₆-alkylsulphonylamino, C₁-C₆alkylsulphonylamino, C₁-C₆alkylsulphonyl

where

R3 and R4 have the meanings indicated above,

or salts, solvates and/or solvates of the salts a salt, solvate or solvate of a salt thereof.

3. (Currently Amended) A compound of Claim claim 1, where

 $R^{1} \hspace*{0.2cm} is \hspace*{0.2cm} C_{1}\text{-}C_{3}\text{-}alkyl \hspace*{0.2cm} or \hspace*{0.2cm} C_{2}\text{-}cycloalkyl, which are optionally substituted by up to 3 } \\ radicals independently of one another selected from the group of $C_{1}\text{-}C_{4}\text{-}alkyl,}$ \\ $C_{1}\text{-}C_{4}\text{-}alkoxy, hydroxycarbonyl, cyano, amino, hydroxy, $C_{1}\text{-}C_{4}\text{-}alkylamino,}$ \\ trifluoromethyl, fluorine, chlorine, bromine, $C_{6}\text{-}C_{10}\text{-}arylcarbonylamino,} $C_{1}\text{-}C_{4}\text{-}alkylaminocarbonyl,} $C_{1}\text{-}C_{4}\text{-}alkylaminocarbonyl,}$ \\ $C_{10}\text{-}arylaminocarbonyl, heteroarylaminocarbonyl,} $C_{1}\text{-}C_{4}\text{-}alkylsulphonylamino,} $$

where C₁-C₄-alkyl and C₁-C₄-alkoxy are optionally substituted by a radical selected from the group of hydroxy, cyano, fluorine, chlorine, bromine, hydroxycarbonyl and a group of the formula –NR³R⁴,

where

R³ and R⁴ are independently hydrogen or C₁-C₄-alkyl,

or

R³ and R⁴ together with the nitrogen atom to which they are bonded are 5- to 6-membered heterocyclyl,

R² is phenyl, pyrimidyl, pyridyl N-oxide or pyridyl, where phenyl is substituted by 1 to 3 radicals and pyrimidyl, pyridyl N-oxide and pyridyl are optionally substituted by 1 to 3 radicals in each case independently of one another selected from the group of C₁-C₄-alkyl, C₁-C₄-alkylamino, hydroxycarbonyl, cyano, trifluoromethyl, amino, hydroxy, C₁-C₄-alkylamino, fluorine, chlorine, bromine, C₆-C₁₀-arylcarbonylamino, C₁-C₄-alkylcarbonylamino, C₁-C₄-alkylcarbonylaminocarbonyl, C₁-C₄-alkylcarbonylaminocarbonyl, heteroarylcarbonylamino, C₁-C₄-alkylsulphonylamino, C₁-C₄-alkylsulphonylamino, C₁-C₄-alkylsulphonyl, and C₁-C₄-alkylthio,

where C₁-C₄-alkyl and C₁-C₄-alkoxy are optionally substituted by a radical selected from the group of hydroxy, cyano, fluorine, chlorine, bromine, hydroxycarbonyl and a group of the formula –NR³R⁴,

where

R3 and R4 have the meanings indicated in Claim claim 1,

or salts, solvates and/or solvates of the salts a salt, solvate or solvate of a salt thereof.

4. (Currently Amended) A compound of Claim claim 1, where

 $R^{1}\,\text{has}$ the meanings indicated in Claim claim 1, and

R² is phenyl, pyridyl N-oxide or pyridyl, where phenyl is substituted by 1 to 3 radicals and pyridyl and pyridyl N-oxide are optionally substituted by 1 to 3 radicals in each case independently of one another selected from the group of

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methyl, ethyl, 2-propyl, trifluoromethyl, methoxy, ethoxy, fluorine and chlorine.

or salts, solvates and/or solvates of the salts a salt, solvate or solvate of a salt thereof.

5. (Currently Amended) A compound Claim of claim 1, where

- R¹ is C₁-C₃-alkyl or C₃-C₆-cycloalkyl, which are optionally substituted by up to 3 radicals independently of one another selected from the group of C₁-C₄-alkyl, trifluoromethyl, fluorine, hydroxy, phenylcarbonylamino, C₁-C₄-alkyl-carbonylamino, C₁-C₄-alkylaminocarbonyl or phenylaminocarbonyl, and
- R² is phenyl, pyridyl N-oxide or pyridyl, where phenyl is substituted by 1 to 3 radicals and pyridyl and pyridyl N-oxide are optionally substituted by 1 to 3 radicals in each case independently of one another selected from the group of methyl, ethyl, 2-propyl, trifluoromethyl, methoxy, ethoxy, fluorine and ehlorine.

or salts, solvates and/or solvates of the salts a salt, solvate or solvate of a salt thereof.

6. (Currently Amended) A compound Claim of claim 1, where

- $$\begin{split} R^1 & \text{is } C_1\text{-}C_3\text{-alkyl or } C_5\text{-}c_9\text{cycloalkyl, which are optionally substituted by up to 3} \\ & \text{radicals independently of one another selected from the group of } C_1\text{-}C_4\text{-alkyl,} \\ & \text{fluorine, trifluoromethyl, hydroxy, phenylcarbonylamino, } C_1\text{-}C_4\text{-alkyl-carbonylamino, } C_1\text{-}C_4\text{-alkyl-aminocarbonyl} \text{ or phenylaminocarbonyl, and} \end{split}$$
- R² is phenyl, pyridyl N-oxide or pyridyl, where phenyl is substituted by one radical and pyridyl and pyridyl N-oxide are optionally substituted by one radical in each case independently of one another selected from the group of methyl, ethyl, 2-propyl, trifluoromethyl, methoxy, ethoxy, fluorine and chlorine.

- (Withdrawn Currently Amended) Process A process for preparing eompounds a compound according to Claim Claim 1, eharacterized in that comprising;
 - [A] converting a compound compounds of the formula

in which

 R^2 has the meanings indicated in Claim claim 1,

are converted by reaction with a compound of the formula

in which R1 has the meanings indicated in Claim claim 1,

and

Z is chlorine or bromine,

in an inert solvent and in the presence of a base, initially into eompounds \underline{a} compound of the formula

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in which

R1 and R2 have the meanings indicated in Claim claim 1,

and then eyelized cyclizing in an inert solvent in the presence of a base to eompounds a compound of the formula (I),

or

[B] <u>reacting a compound</u> empounds of the formula (II) are-reacted with a compound of the formula

in which

R¹ has the meanings indicated in Claim claim 1,

and

R⁵ is methyl or ethyl,

in an inert solvent and in the presence of a base, with direct cyclization to <u>a compound</u> of formula (I),

or

[C] converting a compound compounds of the formula

in which

R² has the meanings indicated in Claim claim 1,

are converted initially by reaction with a compound of the formula (IIIa) in an inert solvent and in the presence of a base into eompounds a compound of the formula

in which

R1 and R2 have the meanings indicated in Claim claim 1,

and the latter are eyelized cyclizing the compound for formula (VI) in a second step in an inert solvent and in the presence of a base and of an oxidizing agent to a compound of (I).

and the resulting compounds of the formula (I) are where appropriate reacted with the appropriate (i) solvents and/or (ii) bases or acids to give their solvates, salts and/or solvates of the salts a salt, solvate or solvate of a salt thereof.

- 8. (Cancelled)
- (Currently Amended) A pharmaceutical composition comprising at least one
 ef the compounds compound of any one of Claims claims 1 to 6 and at least one
 pharmaceutically acceptable, essentially non-toxic carrier or excipient.
- 10. (Cancelled)
- 11. (Cancelled)
- 12. (Cancelled)
- 13. (Withdrawn Currently Amended) A method for the treatment of impairments of perception, concentration, learning and/or memory in humans—or animals a human or animal comprising administering an effective amount of a compound of any one of Claims claims 1 to 6 to the human or animal.

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14. (Withdrawn) The method according to Claim 13, where the impairment is a consequence of Alzheimer's disease.